



SmartPA Criteria Proposal

Drug/Drug Class:	Antihistamines and Antihistamine/Decongestant Combinations, Second Generation PDL Edit
First Implementation Date:	February 26, 2003
Revised Date:	July 7, 2022
Prepared For:	MO HealthNet
Prepared By:	MO HealthNet/Conduent
Criteria Status:	⊠Existing Criteria □ Revision of Existing Criteria
	□New Criteria

Executive Summary

Purpose: The MO HealthNet Pharmacy Program will implement a state-specific preferred drug list.

Why Issue Selected:

The most common form of rhinitis is allergic, occurring in 20 to 40 million patients annually in the US alone. Allergic rhinitis (AR) symptoms range from mild to severe and patients may present with related conditions such as asthma and sinusitis. AR is characterized by nasal mucous membrane swelling and blockage, reflex sneezing, mucous hypersecretion, and often ocular manifestations including itching, tearing, and conjunctival redness. Airborne allergens are known to cause an IgE-mediated response of histamine, thereby beginning the histamine cascade, which creates the role for antihistamine therapy.

Total program savings for the PDL classes will be regularly reviewed.

Program-Specific Information:

fic	Preferred Agents	Non-Preferred Agents
n:	Cetirizine Tabs OTC	Allegra®
	 Cetirizine Soln Rx 	Allegra-D®
	Cetirizine-D	Cetirizine Caps/Chew Tabs/Soln OTC
	 Levocetirizine Tabs Rx 	Clarinex®
	 Loratadine ODT/Soln/Tabs OTC 	Clarinex-D®
	 Loratadine-D 	Desloratadine
		Fexofenadine
		Fexofenadine-D
		Levocetirizine Soln
		Levocetirizine Tabs OTC
		 Loratadine Caps/Chew Tabs OTC
		Quzyttir®
		Zyrtec®

Type of Criteria: ⊠ Increased risk of ADE		☑ Preferred Drug List	
	☐ Appropriate Indications	☐ Clinical Edit	
Data Sources:	☐ Only Administrative Databases	□ Databases + Prescriber-Supplied	

Setting & Population

- Drug class for review: Antihistamines/Decongestant Combinations, Low Sedating (2nd Generation)
- Age range: All appropriate MO HealthNet participants

Approval Criteria

- For Quzyttir: Clinical Consultant Review OR
- For Clarinex syrup: participants aged 2 years or younger AND
- Failure to achieve desired therapeutic outcomes with trial on 2 or more preferred agents
 - Documented trial period of preferred agents OR
 - Documented ADE/ADR to preferred agents

Denial Criteria

- Lack of adequate trial on required preferred agents
- Therapy will be denied if all approval criteria are not met

Claim exceeds maximum dosing limitation for the following:

Drug Description	Generic Equivalent	Max Dosing Limitation
CHILD ZYRTEC 1 MG/ML SOLUTION	CETIRIZINE	10 mL per day
CHILD CETIRIZINE 5 MG CHEW TAB	CETIRIZINE	1 tablet per day
CETIRIZINE HCL 5 MG TABLET	CETIRIZINE	1 tablet per day
CETIRIZINE HCL 5 MG/5 ML SOLN	CETIRIZINE	10 mL per day
ZYRTEC 10 MG CHEWABLE TABLET	CETIRIZINE	1 tablet per day
ZYRTEC 10 MG LIQUID GELS	CETIRIZINE	1 tablet per day
ZYRTEC 10 MG ODT	CETIRIZINE	1 tablet per day
ZYRTEC 10 MG TABLET	CETIRIZINE	1 tablet per day
CLARITIN 10 MG REDITABS	LORATADINE	1 tablet per day
CHILD'S CLARITIN 5 MG TAB CHEW	LORATADINE	1 tablet per day
CLARITIN 10 MG LIQUI-GEL CAP	LORATADINE	1 tablet per day
CLARITIN 5 MG REDITABS	LORATADINE	1 tablet per day
LORATADINE 5 MG/5 ML SOLUTION	LORATADINE	10 mL per day
CLARITIN 5 MG/5 ML SYRUP	LORATADINE	10 mL per day
CLARITIN 10 MG TABLET	LORATADINE	1 tablet per day

	CLARITIN 10 MG TABLET	LORATADINE	1 tablet per day			
Required Documentation						
	boratory Results:	Progress Notes: Other:				
Disposition of Edit						
Denial: Exception Code "0160" (Preferred Drug List) Rule Type: PDL						

Default Approval Period

1 year

References

- Evidence-Based Medicine and Fiscal Analysis: "2nd Generation Antihistamines Therapeutic Class Review", Conduent Business Services, L.L.C., Richmond VA; January 2022.
- Evidence-Based Medicine Analysis: "Non-Sedating Antihistamines", UMKC-DIC; October 2021.
- USPDI, Micromedex; 2022.
- Facts and Comparisons eAnswers (online); 2022 Clinical Drug Information, LLC.